

Complete if Known

Application Number	10/573,890
Filing Date	03-29-06
First Named Inventor	Kazutaka Nakamoto
Art Unit	1625
Examiner Name	Patricia L. Morris
Attorney Docket Number	3939-0118PUS1

(Use as many sheets as necessary)

Sheet	2	of	3
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[illegible][illegible]Date
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3. Office Action that issued the document, by the two-letter code (WIPO Standard ST.3.1.4). For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.7. If possible, 6. Applicant is to place a check mark here if English language Translation is attached.
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STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Sheet 3 of 3

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Group Art Unit	
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NON PATENT LITERATURE DOCUMENTS

Examiner Initial *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	1 2
	23	CHANG et al., "Synthesis and Structure-Activity Relationships of Quaternary Ammonium Cephalosporins with 3-Pyrazolylpyridinium Derivatives," Bioorganic & Medicinal Chemistry Letters (2000) Vol. 10, No. 11, pp. 1211-1214	
	24	CONNORS et al., "Prodrugs in medicine," Overview, Biologicals & Immunologicals, Exp. Opin. Ther. Patents, Vol. 5, No. 9, 1995, pp. 873-885	
	25	Copy of an Office Action from co-pending U.S. Patent Application No. 11/589,128, mailed May 7, 2009	
	26	HATA, "New Approaches to Antifungal Drugs for the Treatment of Fungal and Protozoal Infections, Ravuconazole and Beyond: New Targets and Pre-clinical Strategies," The SMI's 12th Annual Conference, Superbugs and Superdrugs, March 18, 2010, Crowne Plaza London - St. James, 44 pages	
	27	International Search Report dated May 20, 2008 for corresponding International Application No. PCT/JP2008/057851	
	28	ISHIKAWA et al., "TAK-599, a Novel N-Phosphono Type Prodrug of Anti-MRSA Cephalosporin T-91825: Synthesis, Physicochemical and Pharmacological Properties," Bioorganic & Medicinal Chemistry, Vol. 11, pp. 2427-2437, (2003)	
	29	LUKEVICS et al., "Synthesis and cytotoxicity of silyl- and carbonyl-substituted isoxazoles," Chemistry of Heterocyclic Compounds (2000) Vol. 36, No. 10, pp. 1226-1231	
	30	PLATE et al., "Synthesis and Muscarinic Activities of 3-(Pyrazolyl)-1,2,5,6-tetrahydropyridine Derivatives," Bioorganic & Medicinal Chemistry Letters (1996) Vol. 4, No. 2, pp. 227-237	
	31	Supplementary European Search Report dated February 6, 2009 for corresponding European Application No. 04788159.4	
	32	TANAKA et al., "An Effective Lewis Acid-Mediated 1,3-Dipolar Cycloaddition of Nitrile Oxide Using Acetylaldehyde: Synthesis of a (7-Amino-2-pyridyl)isoxazole Derivative and Its Application to Novel Antifungal Agents," pp. 1-8	
	33	VRZHECHCH et al., "Supercooperativity in platelet aggregation: Substituted pyridyl isoxazoles, a new class of supercooperative platelet aggregation inhibitors," FEBS Letters (1994) Vol. 351, No. 2, pp. 168-170	

Examiner
Signature

/Patricia L. Morris/ (07/29/2010)

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* EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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